ElÄºf Uz

# List of Publications by Year in descending order



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|---|---|---|---|
| 14 papers | 827 citations | 840776 11 h-index | 1058476 14 g-index |
| 14 all docs | 14 docs citations | 14 times ranked | 1696 citing authors |

| # | Article | IF | Citations |
|---|---|---|---|
| 1 | Intracerebroventricularly injected nesfatin-1 activates central cyclooxygenase and lipoxygenase pathways. Autonomic Neuroscience: Basic and Clinical, 2020, 226, 102670. | 2.8 | 4 |
| 2 | Loss-of-Function Mutations in ELMO2 Cause Intraosseous Vascular Malformation by Impeding RAC1 Signaling. American Journal of Human Genetics, 2016, 99, 299-317. | 6.2 | 23 |
| 3 | STK4 (MST1) deficiency in two siblings with autoimmune cytopenias: A novel mutation. Clinical Immunology, 2015, 161, 316-323. | 3.2 | 73 |
| 4 | Mutations in the interleukin receptor <scp>IL</scp>11<scp>RA</scp> cause autosomal recessive Crouzonâ€like craniosynostosis. Molecular Genetics & Genomic Medicine, 2013, 1, 223-237. | 1.2 | 70 |
| 5 | Mutations in IRX5 impair craniofacial development and germ cell migration via SDF1. Nature Genetics, 2012, 44, 709-713. | 21.4 | 68 |
| 6 | Disruption of ALX1 Causes Extreme Microphthalmia and Severe Facial Clefting: Expanding the Spectrum of Autosomal-Recessive ALX-Related Frontonasal Dysplasia. American Journal of Human Genetics, 2010, 86, 789-796. | 6.2 | 128 |
| 7 | ALX4 dysfunction disrupts craniofacial and epidermal development. Human Molecular Genetics, 2009, 18, 4357-4366. | 2.9 | 103 |
| 8 | Increased frequency of extremely skewed X chromosome inactivation in juvenile idiopathic arthritis. Arthritis and Rheumatism, 2009, 60, 3410-3412. | 6.7 | 23 |
| 9 | Analysis of skewed X-chromosome inactivation in females with rheumatoid arthritis and autoimmune thyroid diseases. Arthritis Research and Therapy, 2009, 11, R106. | 3.5 | 90 |
| 10 | Skewed X-Chromosome Inactivation in Scleroderma. Clinical Reviews in Allergy and Immunology, 2008, 34, 352-355. | 6.5 | 46 |
| 11 | Mutations in the very low-density lipoprotein receptor VLDLR cause cerebellar hypoplasia and quadrupedal locomotion in humans. Proceedings of the National Academy of Sciences of the United States of America, 2008, 105, 4232-4236. | 7.1 | 88 |
| 12 | Reply to Herz et al. and Humphrey et al. : Genetic heterogeneity of cerebellar hypoplasia with quadrupedal locomotion. Proceedings of the National Academy of Sciences of the United States of America, 2008, 105, E32-3. | 7.1 | 9 |
| 13 | Extremely skewed X-chromosome inactivation is increased in pre-eclampsia. Human Genetics, 2007, 121, 101-105. | 3.8 | 3 |
| 14 | Evidence from autoimmune thyroiditis of skewed X-chromosome inactivation in female predisposition to autoimmunity. European Journal of Human Genetics, 2006, 14, 791-797. | 2.8 | 99 |